Amendments to the Claims

- 1. (original) A chimeric polypeptide which is engineered to include a domain comprising a sequence that directs the attachment of at least one glycosylphosphatidylinositol molecule, wherein said polypeptide is not a ligand binding domain of a cytokine receptor and is for use as a pharmaceutical.
- 2. (currently amended) A <u>The polypeptide</u> according to Claim 1 wherein said polypeptide is a cytokine or variant thereof.
- 3. (currently amended) A <u>The</u> polypeptide according to Claim 1-or 2 wherein said domain comprises the amino acid sequence: PSPTPTETAT PSPTPKPTST PEETEAPSSA TTLISPLSLI VIFISFVLLI (SEQ ID NO: 12).
- 4. (currently amended) A <u>The</u> polypeptide according to Claim 1-or 2 wherein said domain comprises the amino acid sequence:

LVPRGSIEGR GTSITAYNSE GESAEFFFLL ILLLLLVLV (SEQ ID NO: 13).

5. (currently amended) A <u>The</u> polypeptide according to Claim 1 or 2 wherein said domain comprises the amino acid sequence:

TSITAYKSE GESAEFFFLL ILLLLLVLV (SEQ ID NO: 14).

- 6. (currently amended) A <u>The</u> polypeptide according to any of <u>Claims 1-5</u> <u>claim 1</u> wherein said polypeptide includes at least one glycosylphosphatidylinositol molecule.
- 7. (currently amended) A <u>The</u> polypeptide according to <u>any of Claims 2-5 claim 2</u> wherein said polypeptide is selected from the group consisting of: growth hormone; leptin; erythropoietin; prolactin; TNF, interleukins (IL), IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-10, IL-11; the p35 subunit of IL-12, IL-13, IL-15; granulocyte colony stimulating factor (G-CSF); granulocyte macrophage colony stimulating factor (GM-CSF); ciliary neurotrophic factor

Express Mail No. EV629077532US
Date of Deposit: October 7, 2005
Attorney Reference Number 5585-71838-01
Application Number Currently unknown

SLR:dm 10/07/05 426902 P102131US PATENT

(CNTF); cardiotrophin-1 (CT-1); leukemia inhibitory factor (LIF); oncostatin M (OSM); interferon, IFNα and IFNγ,

- 8. (currently amended) A <u>The</u> polypeptide according to any of Claims 1-7 claim 1 wherein said polypeptide has been modified by addition, deletion or substitution of at least one amino acid residue to provide a sequence variant of said polypeptide.
- 9. (currently amended) A <u>The</u> polypeptide according to Claim 8 wherein said variant polypeptide is growth hormone which has been modified in at least one growth hormone receptor binding domain.
- 10. (currently amended) A <u>The</u> polypeptide according to Claim 9 wherein said growth hormone receptor binding domain is in site 1 of growth hormone.
- 11. (currently amended) A The polypeptide according to Claim 9 wherein said growth hormone receptor binding domain is modified in site 2 of growth hormone.
- 12. (currently amended) A The polypeptide according to Claim 9 wherein said growth hormone receptor binding domain is modified in site 1 and site 2 of growth hormone.
- 13. (currently amended) A <u>The</u> polypeptide according to Claim 10-or-12 wherein said modification is selected from the group consisting of: histidine 18 with alanine or aspartic acid; and/or histidine 21 with asparagine; and/or glutamine 22 with alanine; and/or phenylalanine 25 with alanine; and/or aspartic acid 26 with alanine; and/or glutamine 29 with alanine; and/or glutamic acid 167 with alanine; and/or aspartic acid 171 with serine; and/or lysine 172 with serine or alanine; and/or isoleucine 179 with tyrosine, as represented by the growth hormone amino acid sequence in Figure 2 (amino acids 21-254 of SEQ ID NO: 2).
- 14. (currently amended) A <u>The</u> polypeptide according to Claim 13 wherein said modification consists of the the amino acid substitutions: histidine 18 aspartic acid; histidine 21 asparagine;

Express Mail No. EV629077532US
Date of Deposit: October 7, 2005
Attorney Reference Number 5585-71838-01
Application Number Currently unknown

SLR:dm 10/07/05 426902 P102131US PATENT

arginine 167 asparagine; aspartic acid 171 arginine; glutamic acid 174 serine; and isoleucine 179 threonine; as represented by the GH amino acid sequence in Figure 2 (amino acids 21-254 of SEQ ID NO: 2).

- 15. (currently amended) A <u>The</u> polypeptide according to Claim 13 wherein said modification consists of the amino acid substitutions: histidine 18 alanine; glutamine 22 alanine; phenylalanine 25 alanine; aspartic acid 26 alanine; glutamine 29 alanine; glutamic acid 65 alanine; lysine 168 alanine; and glutamic acid 174 alanine; as represented by the GH amino acid sequence in Figure 2 (amino acids 21-254 of SEQ ID NO: 2).
- 16. (currently amended) A <u>The</u> polypeptide according to Claim 11 wherein said site 2 modification is to amino acid residue glycine 120 of the amino acid sequence presented in Figure 2 (amino acids 21-254 of SEQ ID NO: 2).
- 17. (currently amended) A <u>The</u> polypeptide according to Claim 16 wherein said site 2 modification is a substitution of glycine for an amino acid selected from the group consisting of: arginine; alanine; lysine; tryptophan; tyrosine; phenylalanine; and glutamic acid.
- 18. (currently amended) A <u>The</u> polypeptide according to Claim 17 wherein said site 2 substitution is glycine 120 for arginine or lysine or alanine.
- 19. (currently amended) A <u>The</u> polypeptide according to Claim 1 wherein said polypeptide is an antibody.
- 20. (currently amended) A <u>The</u> polypeptide according to Claim 19 wherein said antibody is a monoclonal antibody, or the active binding fragment thereof.
- 21. (currently amended) A <u>The</u> polypeptide according to Claim 20 wherein said monoclonal antibody is a humanised antibody.

Express Mail No. EV629077532US Date of Deposit: October 7, 2005 Attorney Reference Number 5585-71838-01

Application Number Currently unknown

SLR:dm 10/07/05 426902 P102131US PATENT

22. (currently amended) A The polypeptide according to Claim 20 wherein said monoclonal

antibody is a chimeric antibody.

23. (currently amended) A The polypeptide according to Claim 20 wherein the active the

active binding fragment is selected from the group consisting of: F(ab')2, Fab, Fv and Fd

fragments; CDR3 regions; and single chain antibody fragments.

24. (currently amended) A The polypeptide according to Claim 23 wherein said fragment is

a single chain antibody fragment.

25. (currently amended) An oligomeric polypeptide wherein said polypeptide comprises at

least two polypeptides according to any of Claims 1-24 claim 1 which two polypeptides are

linked via a linking molecule.

26. (currently amended) An The oligomeric polypeptide according to Claim 25 wherein said

linker comprises at least one copy of the peptide: Gly Gly Gly Ser (SEQ ID NO: 15).

27. (currently amended) An The oligomeric polypeptide according to Claim 26 wherein said

linker comprises at least 2, 3, 4 or 5 copies of said linker.

28. (currently amended) An The oligomeric polypeptide according to any of Claims 25-27

claim 25 wherein said linker further comprises a protease sensitive cleavage site.

29. (currently amended) An The oligomeric polypeptide according to Claim 28 wherein said

cleavage site is sensitive to a serum protease.

30. (currently amended) An The oligomeric polypeptide according to Claim 29 wherein said

cleavage site comprises the amino acid sequence: LVPRGS (SEQ ID NO: 16).

Page 8 of 11

- 32. (currently amended) An The oligomeric polypeptide according to Claim 29 wherein said cleavage site comprises the amino acid sequence: LVPRGS PGISGGGGGG (SEQ ID NO: 19).
- 33. (currently amended) An The oligomeric polypeptide according to Claim 29 wherein said cleavage site comprises at least two copies of the amino acid sequence SGGGG (SEQ ID NO: 17) which flank said cleavage site.
- 34. (currently amended) An isolated nucleic acid molecule comprising a nucleic acid sequence which encodes a the polypeptide according to claim 1 any of Claims 1-33.
- 35. (currently amended) A vector comprising a the nucleic acid molecule according to Claim
 34.
- 36. (currently amended) A <u>The</u> vector according to Claim 35 wherein said vector is an expression vector adapted for eukaryotic gene expression.
- 37. (currently amended) A cell transfected with the nucleic acid <u>molecule of claim 34 or vector according to any of Claims 34-36</u>.
- 38. (currently amended) A method to prepare the polypeptide of claim 1, a polypeptide or an oligomeric polypeptide according to any of Claims 1-33 comprising:
- i) growing a cell <u>transfected with a nucleic acid sequence which encodes the polypeptide of</u>
 <u>claim 1 according to Claim 37</u> in conditions conducive to the manufacture of said
 polypeptide; and
- ii) purifying said polypeptide from said cell, or its growth environment.

Express Mail No. EV629077532US
Date of Deposit: October 7, 2005
Attorney Reference Number 5585-71838-01
Application Number Currently unknown

SLR:dm 10/07/05 426902 P102131US PATENT

- 39. (currently amended) A cell wherein said cell presents, at least at its cell surface, a polypeptide or oligomeric polypeptide according to <u>claim 1 any of Claims 1-33</u>.
- 40. (currently amended) A method of treatment of an animal, preferably a human, comprising administering an effective amount of a the isolated nucleic acid molecule of claim 34 and/or vector and/or polypeptide and /or cell according to any of Claims 1-37 or 39.
- 41. (new) A method of treatment of an animal, comprising administering an effective amount of the polypeptide of claim 1.
- 42. (new) A method of treatment of an animal, comprising administering an effective amount of the cell of claim 39.